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Bimetallic copper and zinc-catalyzed oxidative cycloaddition of 3-aminopyridazines and nitriles: a direct synthesis of 1,2,4-triazolo[1,5-*b*]pyridazines *via* C–N and N–N bond-forming process†

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One-pot formation of the 1,2,4-triazolo[1,5-*b*]pyridazine nucleus and its derivatives is presented in this manuscript, in which the desired targets are offered easily *via* cooperative Cu(I) and Zn(II)-catalyzed tandem C–N addition and subsequent I₂/KI-mediated intramolecular oxidative N–N bond formation.

The 1,2,4-triazolo[1,5-*b*]pyridazine scaffolds have been recognized an important structural motif in medicinal chemistry, especially showing excellent anti-asthmatic activity.¹ In recent decades, a series of synthetic methods for the synthesis of 1,2,4-triazolo[1,5-*b*]pyridazine derivatives have been elaborated. To date, the preparation of 1,2,4-triazolo[1,5-*b*]pyridazines is typically involved multistep reaction sequences. For example, as shown in Scheme 1, the 3-aminopyridazine derivatives could be converted into *N,N*-dimethylaminomethylene derivatives by treatment with *N,N*-dimethylaminoformamide dimethyl acetal (DMFDMA), and then reacted with hydroxylamine hydrochloride to afford formamide oxime, which treated with polyphosphoric acid to give the fused 1,2,4-triazolo derivatives.^{1,2} In addition, the treatment of 3-aminopyridazine or 3-amino-6-chloropyridazine with *O*-mesitylenesulfonylhydroxylamine (MSH), and the subsequent cyclization with acylating agents, such as formic acid, acetic anhydride and benzoyl chloride, was also an effective procedure to the preparation of 1,2,4-triazolo[1,5-*b*]pyridazine derivatives.³ Although the 1,2,4-triazolo[1,5-*b*]pyridazine derivatives and its analogues could be obtained successfully with multistep reaction sequences in the past decades,⁴ some disadvantages existed in these classic methods, including high temperature, low total yields and highly toxic. Thus the development of one-pot novel methods to access the 1,2,4-triazole nucleus is highly desired.

Recently, one-pot formation of 1,2,4-triazoles nucleus and its analogues using copper-catalyzed tandem addition have also been presented.⁵ And notably, metal-free oxidative N–N bond formation to synthesize 1,2,4-triazoles skeleton was proved to an interesting method for the oxidative construction of heterocycles in the past years.⁶ In comparison to the previously reported multistep methods, these catalytic one-pot transformations provided a simple process for the synthesis of 1,2,4-triazoles. However, to the best of our knowledge, there is no reports on the copper-catalyzed oxidative cycloaddition of 3-aminopyridazines and nitrile for the preparation of 1,2,4-triazolo[1,5-*b*]pyridazine scaffolds. In addition, the development of synthetic methods for the construction of substituted of 1,2,4-triazoles and its derivatives has been of longstanding importance in medicinal chemistry.

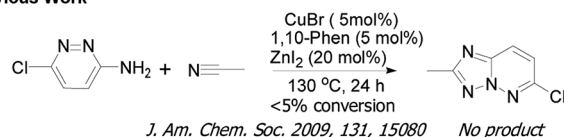
Inspired by previous works on the triazoles,^{5–7} we envisioned a straightforward strategy for the synthesis of the 1,2,4-triazolo[1,5-*b*]pyridazine nucleus by one-pot oxidative cycloaddition reaction of 3-aminopyridazine derivatives and nitriles involving copper-catalyzed C–N bond formation and I₂/KI-mediated

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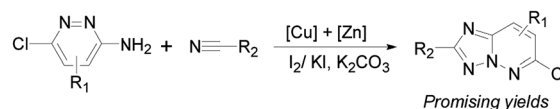
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Previous Work



This Work



Scheme 1 Synthesis of the 1,2,4-triazolo[1,5-*b*]pyridazines.

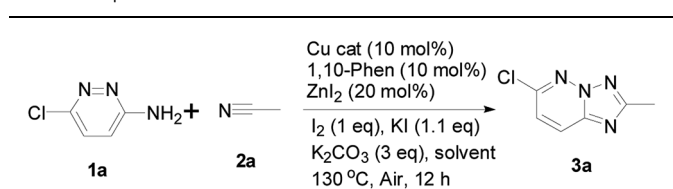
oxidative N–N coupling. Herein we disclose a new and efficient methodology for the preparation of chlorine-containing 1,2,4-triazolo[1,5-*b*]pyridazine scaffolds that difficultly achieved by previous reported methods.

Initially, to explore the feasibility of the copper-based catalytic approach that reported by Ueda and Nagasawa in 2009,⁸ we selected 3-amino-6-chloropyridazine and acetonitrile as a model substrate for reaction evaluation. According to the classic reaction conditions, when **1a** and **2a** were reacted in the presence of 5 mol% of CuBr and 5 mol% of 1,10-phenanthroline as well as 10 mol% ZnI₂ in dichlorobenzene (DCB), at 130 °C for 24 h under air (Scheme 1). Unfortunately, almost no desired product was obtained in this case (Table 1, entry 1). Therefore, the copper-catalyzed oxidative cycloaddition of 3-amino-6-chloropyridazine is different from the copper-catalyzed tandem addition-oxidative cyclization of 2-aminopyridines and nitriles because of its low reactivity under the reported reaction conditions. To explore this approach with copper catalysis, we continued to optimized the cycloaddition reaction of 3-amino-6-chloropyridazine. On the basis of the optimization of reaction conditions, we were pleased to find that the addition of a stoichiometric amount of I₂ or KI to the reaction system is crucial to the formation of desired product **3a** in the presence of K₂CO₃. We hypothesized that the cooperative activation by both CuBr and ZnI₂ was the key step in the addition of 3-amino-6-chloropyridazine to nitriles. To improve the oxidative cycloaddition of 3-amino-6-chloropyridazine to nitrile, we further added I₂ and KI to the reaction system. To our delight, the reaction gave the desired product in promising yield. Similarly to previous report,⁹ I₂/KI-mediated oxidative cyclization is

generally worked *via* the KI₃-promoted N–N bond formation. Notably, no predicted product was observed in the absence of I₂ or K₂CO₃. In addition, the reaction also proceeded in toluene and DMSO, albeit in lower yields. Interestingly, the other copper sources, such as CuI and Cu(OAc)₂, proved to be a negative impact (Table 1, entries 11 and 12).

With the optimized cyclization conditions established above, we further investigated the scope and generality of the synthetic methodology. As shown in Table 2, benzonitriles bearing halogen or trifluoromethyl groups on aromatic rings afforded the corresponding product in moderate yields, while electron-rich benzonitriles reacted with 3-amino-6-chloropyridazine to offer relatively low yields. Delightfully, the heterocyclic nitriles, such as cyanopyridines and cyanothiophene, could also be employed to give moderate yields of products. The reactions performed smoothly when acetonitrile and benzonitriles were employed as substrates to offer desired products in moderate yields. 3-Amino-6-chloropyridazine with electron-donating substituent, such as methoxy group, provided yield slightly higher than that of halogen-containing substrates. In fact, the conversion of 3-aminopyridazines was almost completed, and the starting material was not observed after the reaction. The major reasons included: (1) the desired products were not good in solubility in general organic solvents, thus the corresponding yield was sacrificed during purification by silica gel column chromatography; (2) the side product might be insoluble carbon-based material because this reaction was carried out at high temperature.

Table 1 Optimization of the reaction conditions^a



Entry	Catalyst	Solvent	Additive	Base	Yield ^b (%)
1 ^c	CuBr	DCB	ZnI ₂ /—/—	—	0
2	CuBr	DCB	ZnI ₂ /—/KI	K ₂ CO ₃	0
3	CuBr	DCB	—/I ₂ /KI	K ₂ CO ₃	6
4	CuBr	DCB	ZnI ₂ /I ₂ /—	K ₂ CO ₃	14
5	CuBr	DCB	ZnI ₂ /I ₂ /KI	—	0
6	CuBr	Toluene	ZnI ₂ /I ₂ /KI	K ₂ CO ₃	29
7	CuBr	DMSO	ZnI ₂ /I ₂ /KI	K ₂ CO ₃	24
8	CuBr	DCB	ZnBr ₂ /I ₂ /KI	K ₂ CO ₃	5
9	CuBr	DCB	ZnI₂/I₂/KI	K₂CO₃	40
10	CuBr	DCB	—/I ₂ /—	K ₂ CO ₃	Trace
11	CuI	DCB	ZnI ₂ /I ₂ /KI	K ₂ CO ₃	8
12	Cu(OAc) ₂	DCB	ZnI ₂ /I ₂ /KI	K ₂ CO ₃	Trace

^a Reaction conditions: **1a** (1 mmol, 1 eq.), **2a** (10 mL), CuBr (10 mol%), 1,10-phenanthroline (10 mol%), ZnI₂ (20 mol%), I₂ (1 eq.), KI (1.1 eq.), K₂CO₃ (3 eq.), DCB (2 mL). ^b Determined by LC-MS yield. ^c CuBr (5 mol%), 1,10-phenanthroline (5 mol%), ZnI₂ (10 mol%). DCB is dichlorobenzene.

Table 2 The substrate scope in the Cu/Zn-catalyzed synthesis of the 1,2,4-triazolo[1,5-*b*]pyridazines^a

Entry	R ¹	R ²	Product	Yield ^b (%)	Conversion ^c (%)
1	H	CH ₃	3a	63	>99
2	H	Ph	3b	63	>99
3	H	4-FPh	3c	52	>99
4	H	4-MePh	3d	36	>99
5	H	4-OCF ₃ Ph	3e	18	>99
6	H	2-FPh	3f	34	>99
7	H	2-ClPh	3g	22	>99
8	H	2-OMePh	3h	22	>99
9	H	2-CF ₃ Ph	3i	39	>99
10	H	3-OMePh	3j	27	>99
11	H	2-Thiophen	3k	30	>99
12	H	3-Pyridin	3l	39	>99
13	H	2,4-Cl ₂ Ph	3m	22	>99
14	4-OMe	CH ₃	3n	33	>99

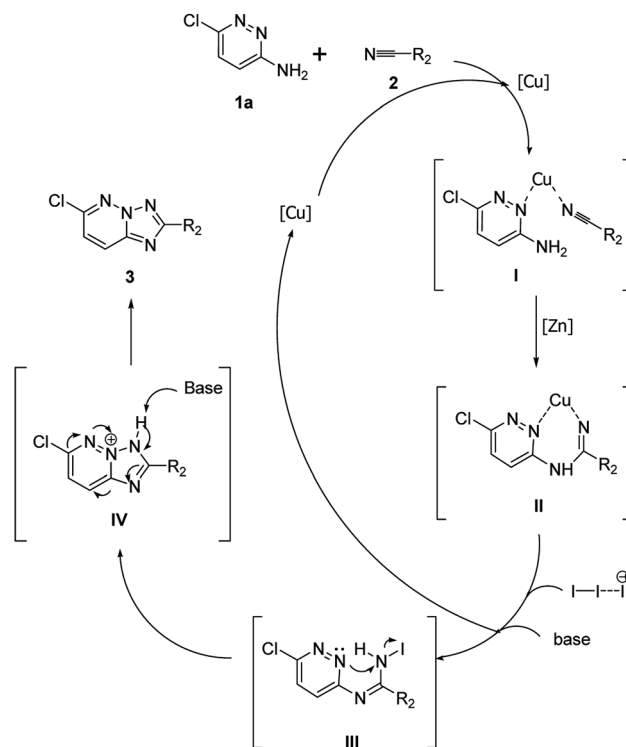
^a Reaction conditions: **1** (0.77 mmol), **2** (1.93 mmol), CuBr (10 mol%), 1,10-phenanthroline (10 mol%), ZnI₂ (20 mol%), I₂ (0.77 mmol), KI (0.85 mmol), K₂CO₃ (2.31 mmol), DCB (5 mL), 130 °C, 12 h, ambient air. ^b Isolated yields. ^c Determined by GC-MS and HPLC.



Notably, the gram-scale synthesis was also carried out. Gratifyingly, the corresponding target **3a** could be obtained in the better yield (75% isolated yield, it is better than that with above small-scale experiment, *versus* 63% yield) under the standard conditions (Scheme 2), which revealed the practical usefulness of present method in the preparation of large amount of target product. In addition, to demonstrate the synthetic utility of this method, **3a** was treated with phenylboronic acid and arylethyne to evaluate its reactivity toward the known Suzuki reaction and Sonogashira reaction. To our delight, the corresponding products were obtained in moderate yields, which may enhance the value of this newly developed method (Scheme 3).

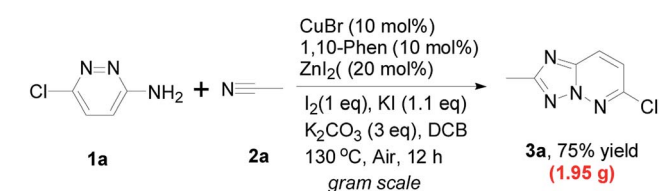
On the basis of the observed results and the LC-MS analysis (see ESI†), we found the side products of this reaction is quite complicated because of the formation of amidine intermediate. Therefore, we suggested a plausible mechanism for the cyclization process of 1,2,4-triazolo[1,5-*b*]pyridazine framework (Scheme 4). The initial amidine intermediate II may be formed firstly *via* copper-catalyzed intramolecular nucleophilic attack of **1a** on the nitrile **2**.¹⁰ Next, amidine intermediate II might provide an iodide intermediate III, promoted by base (K_2CO_3) *via* the subsequent nucleophilic attack of N atom of the pyridazine ring to afford ammonium ion 5, with cleavage of N-I bond.^{6c,9} At the last step, the desired 1,2,4-triazolo[1,5-*b*]pyridazine framework **3** was obtained after the deprotonation and aromatization (Scheme 4).

As described above, there is no reports on the copper-catalyzed oxidative cycloaddition of 3-aminopyridazines and nitrile for the preparation of 1,2,4-triazolo[1,5-*b*]pyridazine scaffolds, which promoted us to consider it as a nitrogen-based

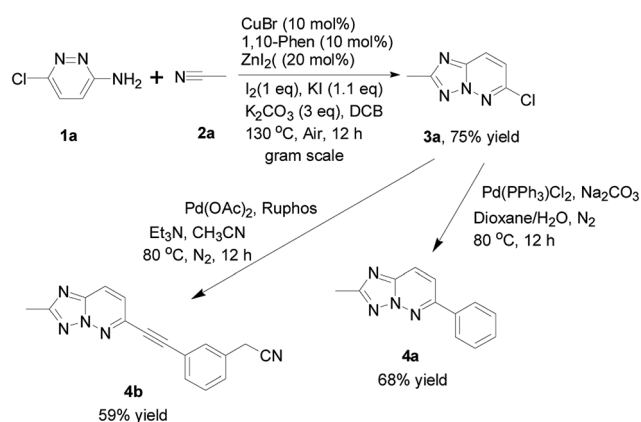


Scheme 4 Propose mechanism of the formation of 1,2,4-triazolo[1,5-*b*]pyridazine **3**.

ligand for copper catalysis because of three nitrogen atoms on this scaffold (Fig. 1). Accordingly, with such a series of 1,2,4-triazolo[1,5-*b*]pyridazine derivatives in hand, we hypothesized that whether the products **3** bearing four nitrogen centers to be worked as ligands in transition metal catalysis. Although no works about 1,2,4-triazolo[1,5-*b*]pyridazine skeleton as ligands was reported in the past, we continued to evaluate its possibility in the copper-catalyzed cycloaddition of carbon dioxide (CO_2) with epoxide. It is well-known that conversion or fixation of CO_2 to synthetically useful compounds has been an important topic for synthetic chemists.¹¹ Among numerous transformations of CO_2 , one of the most attractive and direct synthetic goals starting from carbon dioxide is the construction of five-membered cyclic carbonates because it has widely synthetic uses. And in generally, five-membered cyclic carbonates were achieved from the corresponding diols and phosgene or related compounds.^{11,12} In the past years, we have also demonstrated several catalytic systems for the cycloaddition of epoxides with CO_2 with high efficiency.¹³ However, for most examples in the previous studies, the cycloaddition of CO_2 to epoxides for the preparation of corresponding cyclic carbonates was generally



Scheme 2 Gram-scale synthesis of **3a**.



Scheme 3 Gram-scale reaction and synthetic transformations of **3a**.

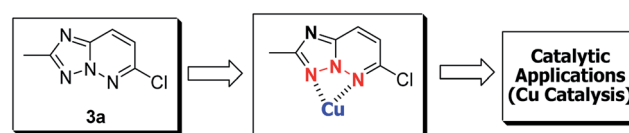


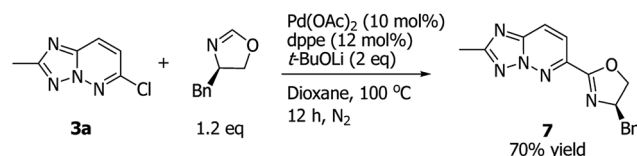
Fig. 1 The structure analysis of 1,2,4-triazolo[1,5-*b*]pyridazine **3a** for copper catalysis.



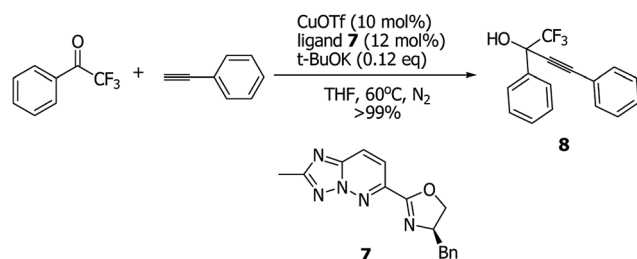
conducted at relatively high reaction temperatures and high pressures (CO₂) in the presence of ionic liquids, metal halides, or metal complexes as catalysts.¹⁴ Therefore, the development of a simple and efficient methodology, especially at atmosphere pressure, for the activation or fixation of CO₂ is a demanding challenge and useful process for organic chemists.

In this part, we would like to applied the cycloaddition of epoxide with carbon dioxide as a model reaction to evaluate the performance of 1,2,4-triazolo[1,5-*b*]pyridazine derivative **3a** as a ligand in the copper catalysis. Initially, many attempts was carried out using **3a** as a ligand, to our delight, the cycloaddition of carbon dioxide with styrene oxide was smoothly took place, and numerous screening experiments about the effect of temperature, pressure of CO₂, as well as the ratio of copper salts with ligand were investigated (Table 3). Under the optimized reaction conditions, the styrene carbonate was obtained in an excellent yield (82%), as shown in Table 3. Therefore present finding provided an alternative method for the copper-catalyzed synthesis of cyclic carbonates from CO₂ and epoxides,¹⁵ which would be an environmentally benign catalyst system in this reaction.

To demonstrate the potential application of 1,2,4-triazolo[1,5-*b*]pyridazine derivative **3a** to the synthesis of other new ligand, the palladium-catalyzed C–H heteroarylation of oxazoline¹⁶ was also utilized as a strategy to give a new 1,2,4-triazolo



Scheme 5 Synthesis of new 1,2,4-triazolo[1,5-*b*]pyridazine-derived oxazoline ligand **7**.



Scheme 6 Copper-catalyzed alkynylation of 2,2,2-trifluoro-1-phenylethanone with 1-ethynylbenzene in the presence of new 1,2,4-triazolo[1,5-*b*]pyridazine-derived oxazoline ligand **7**.

Table 3 Copper-catalyzed cycloaddition of CO₂ with epoxide **5** in the presence of 1,2,4-triazolo[1,5-*b*]pyridazine **3a** as ligand^a

Entry	Cu catalyst	Base	Temp (°C)	Pressure (atm)	Yield ^d (%)
1	CuCl ₂	DMF	100	1	30
2	CuCl ₂	DMAP	100	1	82
3	CuCl ₂	TBAB	100	1	61
4	CuCl ₂	TBAI	100	1	35
5	CuCl ₂	DMAP	70	1	25
6	CuCl ₂	DMAP	130	1	78
7	CuCl ₂	DMAP	160	1	56
8 ^b	CuCl ₂	DMAP	100	1	21
9 ^c	CuCl ₂	DMAP	100	1	52
10	CuBr ₂	DMAP	100	1	78
11	Cu(OAc) ₂	DMAP	100	1	34
12	CuCl ₂	DMAP	100	2	80
13 ^e	CuCl ₂	DMAP	100	1	20
14 ^e	CuCl ₂	—	100	1	NR

^a Note: the ratio of Cu salt with ligand is 1 : 2, except note. ^b With 1 mol% of ligand (**3a**). ^c With 3 mol% of ligand (**3a**). ^d Determined by GC-MS and it is the GC yield. ^e With the ligand **7**.

[1,5-*b*]pyridazine-derived oxazoline ligand **7** (Scheme 5, 70% yield). Catalytic alkynylation of trifluoromethyl ketones is an important reaction in organic synthesis and organofluorine chemistry, which provided a facile process to the preparation of fluorinated propargylic alcohols.¹⁷ And in the past years, many methods have been reported for the catalytic synthesis of such fluorinated alcohols by catalytic alkynylation of trifluoromethyl ketones.¹⁸ However, development of highly efficient benign methodologies for alkynylation of trifluoromethyl ketones is still very important to the advancement of synthetic organic chemistry. In this work, under the Shibasaki's reported reaction conditions,^{18b} we found the new 1,2,4-triazolo[1,5-*b*]pyridazine-derived oxazoline **7** was a highly efficient ligand in this reaction (>99% yield, see Scheme 6), albeit without enantioselectivity in this case. We believed that the described reactions in this work could be extended to the synthesis of structurally diverse propargyl alcohols.

In summary, we have presented a novel and simple metal-catalyzed tandem oxidative cycloaddition reaction, in which the desired targets are offered easily *via* cooperative Cu(I) and Zn(II)-catalyzed tandem C–N addition and subsequent I₂/KI-mediated intramolecular oxidative N–N bond formation. Although the yields of the desired products were not perfect at present, the method will be an attractive alternative for the preparation of potentially biological active 1,2,4-triazolo[1,5-*b*]pyridazine derivatives, with features such as the broad substrate scope within short reaction time. The future applications of this compound class and its derivatives in organic synthesis, medicinal chemistry, and pesticide chemistry are still ongoing in our laboratory.

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Notes and references

- (a) M. Kuwahara, Y. Kawano, M. Kajino, Y. Ashida and A. Miyake, *Chem. Pharm. Bull.*, 1997, **45**, 1447; (b) M. Gyoten, H. Nagaya, S. Fukuda, Y. Ashida and Y. Kawano, *Chem. Pharm. Bull.*, 2003, **51**, 122; (c) S. D. Edmondson, A. Mastracchio, R. J. Mathvink, J. He, B. Harper, Y. J. Park, M. Beconi, J. D. Salvo, G. J. Eiermann, H. He, B. Leitling, J. F. Leone, D. A. Levorse, K. Lyons, R. A. Patel, S. B. Patel, A. Petrov, G. Scapin, J. Shang, R. S. Roy, A. Smith, J. K. Wu, S. Xu, B. Zhu, N. A. Thornberry and A. E. Weber, *J. Med. Chem.*, 2006, **49**, 3614; (d) C. J. Menet, S. R. Fletcher, G. V. Lommen, R. Geney, J. Blanc, K. Smits, N. Jouannigot, P. Deprez, E. M. van der Aar, P. Clement-Lacroix, L. Lepescheux, R. Galien, B. Vayssiere, L. Nelles, T. Christophe, R. Brys, M. Uhring, F. Ciesielski and L. V. Rompaey, *J. Med. Chem.*, 2014, **57**, 9323.
- (a) S. Polanc, B. Vercek, B. Stanovnik and M. Tisler, *Tetrahedron Lett.*, 1972, 1677; (b) S. Polanc, B. Vercek, B. Stanovnik and M. Tisler, *J. Org. Chem.*, 1974, **39**, 2143; (c) B. Stanovnik, A. Stimac, M. Tisler and B. Vercek, *J. Heterocycl. Chem.*, 1982, **19**, 577; (d) B. Stanovnik, V. Stibilj and M. Tisler, *Synthesis*, 1986, 807.
- Y. Tamura, J. H. Kim and M. Ikeda, *J. Heterocycl. Chem.*, 1975, **12**, 107.
- (a) M. Zupan, B. Stanovnik and M. Tisler, *Tetrahedron Lett.*, 1972, 4179; (b) G. Schneider and M. Nettekoven, *J. Comb. Chem.*, 2003, **5**, 233; (c) D. G. Yu, M. Suri and F. Glorius, *J. Am. Chem. Soc.*, 2013, **135**, 8802; (d) K. L. Stevens, M. J. Reno, J. B. Alberti, D. J. Price, L. S. Kane-Carson, V. B. Knick, L. M. A. M. Hassell, J. M. Veal, S. T. Davis, R. J. Griffin and M. R. Peel, *Bioorg. Med. Chem. Lett.*, 2008, **18**, 5758; (e) W. M. Abdou, N. A. Ganoub and E. Sabry, *Beilstein J. Org. Chem.*, 2013, **9**, 1730; (f) T. Irrgan and R. Kempe, *Eur. J. Org. Chem.*, 2005, 4382.
- (a) J.-P. Zhang, Y. Y. Lin, X. C. Huang and X. M. Chen, *J. Am. Chem. Soc.*, 2005, **127**, 5495; (b) X. Meng, C. Yu and P. Zhao, *RSC Adv.*, 2014, **4**, 8612; (c) B. Bartels, C. G. Bolas, P. Cueni, S. Fantasia, N. Gaeng and A. S. Trita, *J. Org. Chem.*, 2015, **80**, 1249; For recent reviews on copper-catalyzed organic reactions, see: (d) A. E. Allen, R. R. Walvoord, R. Padilla-Salinas and M. C. Kozlowski, *Chem. Rev.*, 2013, **113**, 6234.
- (a) O. Prakash, H. K. Gujral, N. Rani and S. P. Sing, *Synth. Commun.*, 2000, **30**, 417; (b) Z. Zheng, S. Ma, L. Tang, D. Z. Negrierie, Y. Du and K. Zhao, *J. Org. Chem.*, 2014, **79**, 4686; (c) L. Song, X. Tian, Z. Lv, E. Li, J. Wu, Y. Liu, W. Yu and J. Chang, *J. Org. Chem.*, 2015, **80**, 7219.
- (a) Z. J. Zheng, F. Ye, L. S. Zheng, K. F. Yang, G. Q. Lai and L. W. Xu, *Chem.-Eur. J.*, 2012, **18**, 14094; (b) C. Y. Wang, J. F. Zou, Z. J. Zheng, W. S. Huang, L. Li and L. W. Xu, *RSC Adv.*, 2014, **4**, 54256; (c) T. Song, L. Li, W. Zhou, Z. J. Zheng, Y. Deng, Z. Xu and L. W. Xu, *Chem.-Eur. J.*, 2015, **21**, 554; (d) M. Y. Chen, T. Song, Z. J. Zheng, Z. Xu, Y. M. Cui and L. W. Xu, *RSC Adv.*, 2016, **6**, 58698.
- S. Ueda and H. Nagasawa, *J. Am. Chem. Soc.*, 2009, **131**, 15080.
- (a) P. H. Svensson and L. Kloo, *Chem. Rev.*, 2003, **103**, 1650; (b) P. Gogoi and D. Konwar, *Org. Biomol. Chem.*, 2005, **3**, 3473; (c) X. Tian, L. Song, M. Wang, Z. Lv, J. Wu, W. Yu and J. Chang, *Chem.-Eur. J.*, 2016, **22**, 7617.
- (a) G. Rousselet, P. Capdevielle and M. Maumy, *Tetrahedron Lett.*, 1993, **34**, 6395; (b) W. Yin, C. Wang and Y. Huang, *Org. Lett.*, 2013, **15**, 1850; (c) Z. Zheng, S. Ma, L. Tang, D. Zhang-Negrierie, Y. Du and K. Zhao, *J. Org. Chem.*, 2014, **79**, 4687; (d) H. Xu, S. Ma, Y. Xu, L. Bian, T. Ding, X. Fang, W. Zhang and Y. Ren, *J. Org. Chem.*, 2015, **80**, 1789; (e) C.-y. Chen, G. Tang, F. He, Z. Wang, H. Jing and R. Faessler, *Org. Lett.*, 2016, **18**, 1690.
- For representative reviews, see: (a) D. H. Dibson, *Chem. Rev.*, 1996, **96**, 2063; (b) X. B. Lu and D. J. Darensbourg, *Chem. Soc. Rev.*, 2012, **41**, 1462; (c) A. M. Appel, J. E. Bercaw, A. B. Bocarsly, H. Dobbek, D. L. Dubois, M. DuPuis, J. G. Ferry, E. Fujita, R. Hille and P. J. A. Kenis, *Chem. Rev.*, 2013, **113**, 6621; (d) J. F. Shi, Y. J. Jiang, Z. Y. Jiang, X. Y. Wang, X. L. Wang, S. H. Zhang, P. P. Han and C. Yang, *Chem. Soc. Rev.*, 2015, **44**, 5981; (e) H. Wang, J. J. Peng and J. Li, *Chem. Rec.*, 2016, **16**, 1298; (f) L. J. Guo, Y. J. Wang and T. He, *Chem. Rec.*, 2016, **16**, 1918; (g) J. Hasegawa, R. Miyazaki, C. Maeda and T. Ema, *Chem. Rec.*, 2016, **16**, 2260; For recent examples, see: (h) G. P. Ji, Z. Z. Yang, H. Y. Zhang, Y. F. Zhao, B. Yu, Z. S. Ma and Z. M. Liu, *Angew. Chem., Int. Ed.*, 2016, **55**, 9684; (i) W. Y. Gao, H. F. Wu, K. Y. Leng, Y. Y. Sun and S. Q. Ma, *Angew. Chem., Int. Ed.*, 2016, **55**, 5472; (j) J. Rintjema, R. Epping, G. Fiorani, E. Martin, E. C. Escudero-Adan and A. W. Kleij, *Angew. Chem., Int. Ed.*, 2016, **55**, 3972; (k) D. Y. Zhang, S. K. Boopathi, N. Hadjichristidia, Y. Gnanou and X. S. Feng, *J. Am. Chem. Soc.*, 2016, **138**, 11117; (l) C. Romain, Y. Q. Zhu, P. Dingwall, S. Paul, H. S. Rzepa, A. Buchard and C. K. Williams, *J. Am. Chem. Soc.*, 2016, **138**, 4120.
- (a) V. Amarnath and A. D. Broom, *Chem. Rev.*, 1977, **77**, 183; (b) M. Pena-Lopez, H. Neumann and M. Beller, *Eur. J. Org. Chem.*, 2016, 3721; (c) G. L. Gregory, M. Ulmann and A. Buchard, *RSC Adv.*, 2015, **5**, 39404; (d) F. D. Bobbink, W. Gruszka, M. Hulla, S. Das and P. J. Dyson, *Chem. Commun.*, 2016, **52**, 10787; (e) S. B. Wang and X. C. Wang, *Angew. Chem., Int. Ed.*, 2016, **55**, 2308.
- (a) F. W. Li, C. G. Xia, L. W. Xu, C. G. Xia, W. Sun and G. X. Chen, *Chem. Commun.*, 2003, 2042; (b) F. W. Li, L. F. Xiao, C. G. Xia and B. Hu, *Tetrahedron Lett.*, 2004, **45**, 8307; (c) L. W. Xu, M. S. Yang, J. X. Jiang, H. Y. Qiu and G. Q. Lai, *Cent. Eur. J. Chem.*, 2007, **5**, 1073; (d) F. Wang, C. Z. Xu, Z. Li, C. G. Xia and J. Chen, *J. Mol. Catal. A: Chem.*, 2014, **385**, 133; (e) H. L. Liu, Z. Huang, Z. Han, K. L. Ding, H. C. Liu, C. G. Xia and J. Chen, *Green Chem.*, 2015, **17**, 4281.



- 14 (a) M. North, R. Pasquale and C. Young, *Green Chem.*, 2010, **12**, 1514; (b) T. Sakakura and K. Kohno, *Chem. Commun.*, 2009, 1312; (c) A. Decortes, A. M. Castilla and A. W. Kleij, *Angew. Chem., Int. Ed.*, 2010, **49**, 9822; (d) M. Cokoja, C. Bruckmeier, B. Rieger, W. A. Herrmann and F. E. Kühn, *Angew. Chem., Int. Ed.*, 2011, **50**, 8510.
- 15 (a) L. Wu, H. Yang, H. Wang and J. Lu, *RSC Adv.*, 2015, **5**, 23189; (b) M. Y. Wang, Q. W. Song, R. Ma, J. N. Xie and L. N. He, *Green Chem.*, 2016, **18**, 282.
- 16 T. Xi, Y. Mei and Z. Lu, *Org. Lett.*, 2015, **17**, 5939.
- 17 (a) C. J. Li, *Acc. Chem. Res.*, 2010, **43**, 581; (b) J. Nie, H. C. Guo, D. Cahard and J. A. Ma, *Chem. Rev.*, 2011, **111**, 455.
- 18 (a) R. Motoki, D. Tomita, M. Kanai and M. Shibasaki, *Tetrahedron Lett.*, 2006, **47**, 8083; (b) R. Motoki, M. Kanai and M. Shibasaki, *Org. Lett.*, 2007, **9**, 2997; (c) G. J. Deng and C. J. Li, *Synlett*, 2008, 1571; (d) G. W. Zhang, W. Meng, H. Ma, J. Nie, W. Q. Zhang and J. A. Ma, *Angew. Chem., Int. Ed.*, 2011, **50**, 3538–3542; (e) V. R. Chintareddy, K. Wadhwa and J. G. Verkade, *J. Org. Chem.*, 2011, **76**, 4482; (f) C. A. Correia, D. T. McQuade and P. H. Seeberger, *Adv. Synth. Catal.*, 2011, **355**, 3517; (g) H. Wang, K. F. Yang, L. Li, Y. Bai, Z. J. Zheng, W. Q. Zhang, Z. W. Gao and L. W. Xu, *ChemCatChem*, 2014, **6**, 580; (h) L. Wang, N. Liu, B. Dai, X. Ma and L. Shi, *RSC Adv.*, 2015, **5**, 10089; (i) F. Lazreg, M. Lesieur, A. J. Samson and C. S. J. Cazin, *ChemCatChem*, 2016, **8**, 209; (j) P. Czerwinski, E. Molga, L. Cavallo, A. Poater and M. Michalak, *Chem.–Eur. J.*, 2016, **22**, 8089; (k) J.-i. Ito, S. Ubukata, S. Muraoka and H. Nishiyama, *Chem.–Eur. J.*, 2016, **22**, 16801; (l) Y. Zheng, Y. Tan, K. Harms, M. Marsch, R. Riedel, L. Zhang and E. Meggers, *J. Am. Chem. Soc.*, 2017, **139**, 4322; (m) F. L. Li, L. Wang, C. H. Li, N. Liu and B. Dai, *ACS Omega*, 2017, **2**, 1104; (n) X. D. Li, H. Y. Ma, C. H. Xing and L. Lu, *Tetrahedron Lett.*, 2017, **58**, 1564.

